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An ethnobotanical survey and antifungal activity of *Piper guineense* used for the treatment of fungal infections in West-African traditional medicine

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Abstract

Ethnobotanical relevance: *Piper guineense* occurs commonly in West Africa where it is used for fungal infections instead of the costly and not always accessible conventional antifungals. Fungal, yeast-based diseases are common in West-Africa especially among those living with HIV/ AIDS, and thus this study was performed in Imo state, South-Eastern Nigeria, where *P. guineense* is predominantly used for the treatment of fungal diseases, such as skin rashes, oral thrush and vaginosis.

Aim of study: The scarce number of previous studies on the documentation of the traditional uses of *P. guineense* extracts for the treatment of fungal infections in Nigeria prompted this survey. The investigation focused on how traditional healers recognize and diagnose fungal infections, how *P. guineense* is collected, on the various parts used for the treatments, methods of preparations, administrations and treatments. In addition, an *in vitro* antifungal screening of *P. guineense* fruit and leaf extracts of various polarities, and piperine and piperlongumine, representing the main constituents in these extracts, were performed.

Methods: A house to house ethnobotanical survey was conducted using questionnaires. Twenty traditional medical practitioners (TMP) and herb sellers from ten villages were interviewed. Four human pathogenic strains of yeast and *Cryptococcus neoformans*, a yeast-like basidiomycete causing meningitis in immunocompromised individuals, were used for the antifungal screening.

Results: The traditional medical practitioners (TMP) and herb sellers explained that the leaves and fruits are the most commonly used plant parts for the treatments. The oral intake of the extracts in locally produced bamboo alcohol (Kai-kai) is the most common method of administration. In

accordance with these recorded traditional uses, we found that extracts of *P. guineense* were growth inhibitory against the fungal strains with MIC values ranging from 39 to 2500 µg/mL. The lowest MIC value of 39 µg/mL was recorded for a methanol fruit extract against *Candida albicans*, *C. glabrata* and *C. tropicalis*. In addition, ethanol and hexane fruit extracts were effective against the growth of *C. albicans* and *C. glabrata*, respectively, with a MIC of 78 µg/mL. Piperlongumine and piperine were active against *C. albicans* with MIC values of 39 and 78 µg/mL respectively.

Conclusion: *P. guineense* fruit and leaf extracts, as well as their piperamide alkaloid constituents piperine and piperlongumine, have interesting antifungal properties and could have potential as new antifungal scaffolds. Our results warrant further in-depth investigations to isolate and characterize piperamide alkaloids and other compounds responsible for the antifungal activity in the extracts.

Key words: *Piper guineense*, antifungal uses, ethnobotanical survey, antifungal screening, piperine, piperlongumine

Introduction

Africa is richly endowed with medicinal plants that are utilized in the treatment of fungal diseases in traditional medicine and thus could be sources for standardized antifungal extracts and antifungal compounds. Examples of such plant species are *Jatropha curcas* L. (Euphorbiaceae), *Monodora myristica* Dunal. (Annonaceae), *Xylopia aethiopica* A. Rich. (Annonaceae) and *Piper guineense* Schumacher & Thonn (Piperaceae) (Ngane et al., 2003; Abdelgadir and Van Staden, 2013; Dzoyem and Kuete, 2013). The mortality rate associated with fungal diseases is high in Africa especially among those living with HIV/ AIDS (Park et al., 2009). Due to the high cost of conventional medicine and accessibility of numerous medicinal plants in Sub-Saharan Africa, 70- 80% of the population use traditional medicine and plant-based preparations for the treatment of infectious diseases (Agyare et al., 2018). In most rural villages in Africa, the conventional health-care professionals are limited and there are no adequate health-care facilities, and thus, majority of the people patronize the traditional medical practitioners who use the readily available herbal formulations for the treatment of fungal infections (Osuchukwu et al., 2017).

African guinea pepper (*Piper guineense* Schumacher & Thonn) is a medicinal plant and a spice that is highly valued in Africa because of its numerous ethno-medicinal uses (Besong et al., 2016; Ajibesin et al., 2011, Mgbeahuruiké et al., 2017, 2018). The extracts of *P. guineense* are used to treat topical fungal diseases in West-African traditional medicine (Ngane et al., 2003). It is a vine that can grow up to 20 m in length with edible leaves and peppercorn fruits that have a strong pungent aroma

(Freiesleben et al., 2015). *P. guineense* is used for the preparation of herbal formulations for the treatment of numerous ailments and infectious diseases (Obodozie et al., 2010; Asase et al., 2012). Research has shown that fractions, natural compounds and extracts from *P. guineense* can be explored as antifungal preparations and agents for the prevention of skin infections (Ngane et al., 2003). In another study, it was established that *P. guineense* extracts could be used as botanical fungicides by local farmers because of the high cost and toxicity associated with synthetic fungicides (Asawalam, 2006). According to previous research, herbal formulations from medicinal plants are often more effective and safer than those from isolated active compounds (Fabricant and Farnsworth, 2001), and thus standardized extracts could be used in the combat against fungal infections to minimize the risk of the development of resistance.

Table 1. Summary of ethno-medicinal uses, pharmacology/antifungal screening of *Piper guineense*.

<i>P. guineense</i> plant part	Ethno-medicinal uses with reference source	Pharmacology/ Antifungal activity with reference source	<i>In vivo/ In vitro</i>
Seeds/ Fruits	Seeds are used to prepare decoctions for the treatment of fungal infections (Ngane et al., 2003). Seeds and fruits in combination with other medicinal plants such as <i>Xylopiya aethiopica</i> (Annonaceae) is boiled and administered to nursing mothers to aid the contraction of the uterus (Okigbo and Igwe, 2007).	Agar diffusion method with 90% ethanolic extract and fractions obtained by column chromatography. Amphotericin B, Griseofulvin and Clotrimazole at 10 µg/mL each were used as positive controls. MIC against <i>Candida albicans</i> , <i>Cryptococcus neoformans</i> , <i>Scopulariopsis brevicaulis</i> , <i>Aspergillus flavus</i> , and <i>Microsporum gypseum</i> were within the range of 50 -100 µg/mL for the extracts (Ngane et al., 2003).	<i>In vitro</i> and <i>in vivo</i> (Ngane et al., 2003).
	Seeds are used to prepare decoctions for the treatment of asthma and psychotic disorder (Gbekley et al., 2017; Oyemitan et al., 2015). The fruits are used for respiratory infections, female infertility, and aphrodisiac (Dzoyem et al., 2014). Fruits are used to prepare decoctions with other herbs to treat epilepsy (Abila et al. 1993).	The fruits were extracted with a mixture of methanol-dichloromethane (3:1 V/V) for 48 h at room temperature. Broth micro-dilution method was used, Nystatin was used as positive control, MIC was between 1.56 to 6.25 mg/mL against <i>Candida albicans</i> , <i>Candida parapsilosis</i> , <i>Candida tropicalis</i> , <i>Candida krusei</i> , <i>Candida glabrata</i> and <i>Cryptococcus neoformans</i> (Dzoyem et al., 2014)	<i>In vitro</i> (Dzoyem et al., 2014).
	Used for cough, bronchitis, rheumatism and syphilis (Konning et al., 2004; Sandberge et al., 2005). The fruits are used to prepare Yaji soup which is eaten as an aphrodisiac in West African countries (Ibrahim et al., 2010). It is mixed with other herbs to produce Niprisan used in herbal medicine in West Africa for the treatment of sickle cell anemia (Obodozie et al., 2010, Freiesleben et al., 2015).	Agar diffusion method, Soxhlet 96% methanol extracts. Clotrimazole was used as a positive control, inhibition zones were between 7.4 to 9.8 mm against <i>Candida albicans</i> and <i>Aspergillus niger</i> (Konning et al., 2004).	<i>In vitro</i> (Konning et al., 2004).
		Petroleum ether extract obtained by Soxhlet extraction, aqueous extract and 95 % ethanolic extract. MIC was 200 mg/mL against <i>Aspergillus niger</i> , No positive control was mentioned (Ebana et al., 2016).	<i>In vitro</i> (Ebana et al., 2016).
Leaves	Leaves have anticonvulsive	Petroleum ether extract obtained by	<i>In vitro</i> (Ebana et al., 2016).

	activities and are used to make decoctions for the treatment of convulsion (Abila et al., 1993). Leaves are boiled with <i>Xylopia aethiopica</i> and taken for the treatment of malaria (Asase et al., 2012). Leaves are used to prepare soup for the treatment of sexually transmitted diseases (Ajibesin et al., 2011). Used for the treatment of dysentery and bronchitis (Ogunniran et al., 2009).	Soxhlet extraction, aqueous extract and 95 % ethanolic extract. MIC was 80 mg/mL against <i>Aspergillus niger</i> , Significant antifungal activity was observed. No positive control was mentioned (Ebana et al., 2016). Hexane and ethanol extracts. The diameter of the inhibition zones were 11.0-34.0 mm against <i>Aspergillus niger</i> and <i>Aspergillus flavus</i> (Dada et al., 2013). Water extracts obtained by maceration. 5, 10 and 15 % concentration of the water extract was used. The extracts were effective against <i>Fusarium oxysporum</i> at higher concentrations of 10 and 15 % (Abiala et al., 2015).	<i>In vitro</i> (Dada et al., 2013). <i>In vitro</i> (Abiala et al., 2015).
Root	No references found	No references found	No references found

80

81 This present study is the first report resulting from an ethnobotanical survey on the traditional uses
82 of *P. guineense* extracts for the treatment of fungal infections, including candidiasis, in South-
83 Eastern Nigeria. Due to the frequent occurrence of fungal diseases in West Africa, this study was
84 performed in Imo state, South-Eastern Nigeria, where *P. guineense* is predominantly used for the
85 treatment of fungal diseases and for other various medicinal purposes. Imo state is one of the five
86 states in South-Eastern Nigeria. The people in Imo state belong to the Igbo ethnic group and are
87 predominantly Christians. This study area was selected because of the prevalence of fungal
88 infections in the area, owing to the fact that it is a rain forest zone and the humid climate favours the
89 occurrence of topical fungal infections. The area was also selected due to the fact that no previous
90 investigations on the ethnopharmacology of *P. guineense* have been performed in this region.
91 Moreover, the area is inhabited by over 4.8 million people, who largely cultivate *P. guineense* and
92 depend on this species and other medicinal plants for the treatment of infectious diseases and
93 among them fungal infections. Also, the people have a diverse cultural and historical background,
94 and are well grounded with the knowledge of traditional medicine. However, this valuable
95 traditional medicinal knowledge is usually not documented. Thus, the aim of the study was to
96 conduct an ethnobotanical survey in order to document the ways in which traditional healers and
97 herb sellers utilize *P. guineense* extracts and preparations for the treatment of fungal diseases. The
98 investigation was focused on how fungal infections are recognized and diagnosed, how and which
99 parts of *P. guineense* are collected, methods of preparations, administrations and treatments. To
100 validate the ethnobotanical information, an *in vitro* screening of *P. guineense* extracts of the fruit and
101 leaf was performed against selected human pathogenic *Candida* spp. and *Cryptococcus neoformans*
102 in search for new antifungal extracts and agents.

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2. Materials and methods

2.1 Study area and survey

Imo state is situated in South-Eastern Nigeria and share boundary with the South-South geo-political zone of Nigeria. It is a tropical rain forest zone located between 4°45 -7°15N and 6°50 - 7°25E, covering a total land area of 5,530 km² (Fig. 1). An ethnobotanical survey was conducted in Imo state, where *P. guineense* is frequently used as herbal remedy in the treatment of candidiasis and other fungal diseases. The field work was conducted in ten localities between November and December 2017. A total of 20 traditional healers and herb sellers were interviewed (two from each locality). The purpose of the work was to validate the use of *P. guineense* in the treatment of fungal diseases. The 10 localities visited during the interviews were chosen by random sampling technique and are not clustered, but are all distributed within the three senatorial districts in Imo state as seen in Figure 1. The two traditional healers interviewed from each locality were independent from each other.

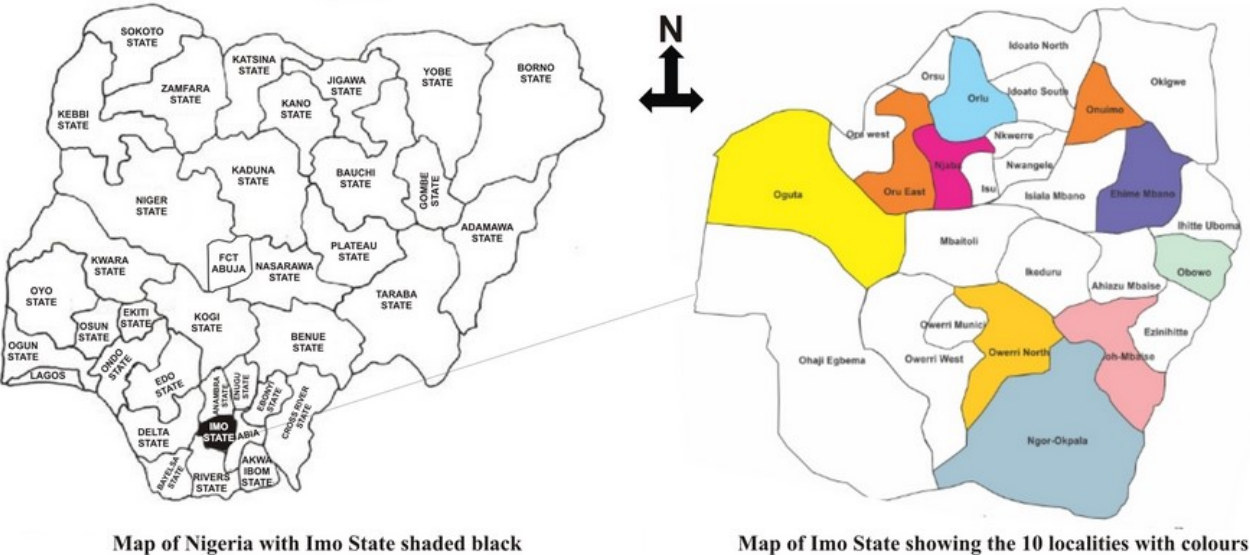


Fig 1. Map of Nigeria showing the study area (Imo state) shaded in black and Map of Imo state showing the 10 localities shaded in colours. Source, Eunice Ego Mgbeahuruike, University of Helsinki, Finland and Chijioke Chukwueke (Graphic designer).

126 2.2 Data collection

127 In the ethnobotanical survey, a house to house strategy was used with the permission of the village
128 heads and traditional rulers. First, an ethical consideration of the project was done appropriately and
129 approved by the University of Helsinki ethical committee, the survey plan was also approved by the
130 head of the Division of Pharmaceutical Biosciences of the University of Helsinki, Finland. The
131 village heads were notified prior to the investigation and the traditional healers were also well
132 informed. The investigating team obtained an ethical approval from each of the village heads. The
133 individual traditional healers gave an informed consent for the use of the data and pictures obtained
134 during the interview. The consent was an oral agreement which was obtained from each of the
135 traditional healers before the interviews began and permission to publish any photographs resulting
136 from the interviews was obtained too. Detailed and validated questionnaires (Appendix A) were
137 administered to twenty traditional healers (2 from each locality). This was followed by series of
138 questions and conversations, so as to obtain the information needed for the work. The
139 questionnaires were written in English, but the local language (Igbo) was used during the interviews
140 and conversations because most of the participants could not interact well in English even though
141 they had formal education. The traditional healers provided samples of the leaves, and fruits of *P.*
142 *guineense* which was identified by a member of the investigating team who is a botanist. The
143 traditional healers cooperated with our investigating team and some cash gifts were given to them at
144 the end of the survey.

145

146 2.3 Plant material for antifungal screening

147 The fruits and leaves of *P. guineense* (African guinea pepper) used in this study were collected from
148 a rural village in Imo State, South Eastern Nigeria. The plant materials were authenticated at the
149 Department of Crop Science of the Federal University of Technology, Owerri, Nigeria. Voucher
150 specimens are deposited in the herbarium of the Department of Crop Science of the same university
151 with the specimen number FUTO/SAAT/NS/005A for the fruit and FUTO/SAAT/NS/005B for the
152 leaf.

153

154 2.4 Extraction

155 The air dried plant materials were milled with a grinder to obtain finely ground powdery samples.
156 Sequential extraction was carried out using solvents of varying polarities, starting with the least
157 polar solvent. First, 40 g of the plant material was extracted with 300 mL of hexane, followed by
158 extraction with 300 mL of chloroform, then 300 mL of ethanol and the residue was finally extracted
159 and washed with 300 mL of methanol. Fruits and leaf plant materials were used for the extractions.

160 The extraction with each of the solvents was conducted in duplicates and each extract, except from
161 the water extracts, were filtered using filter paper (Whatman GE Healthcare, Chicago, IL, USA).
162 The water extracts were centrifuged at 689 g for 5 minutes. For each extract, the filtrate was
163 evaporated using a rotary evaporator (Rotavapor, Heidolph VV2000) combined with a water bath
164 not exceeding +40°C, thereafter the extracts were lyophilized for two days to dry completely.
165 Macerations and hot water decoctions were also prepared from the plant samples since these
166 preparations are used in traditional medicine. Macerations were prepared by weighing 10 g of the
167 fruits and leaf plant materials into Erlenmeyer flasks. 100 mL of water was added and extraction
168 was performed for 24 hours using a magnetic stirrer. The mixture was centrifuged at 689 g for 15
169 min (Eppendorf AG centrifuge 5810R, Germany). For the decoctions, 10 g of the plant material was
170 boiled with 100 mL of water and allowed to cool. The mixture was centrifuged for 15 min at 689 g
171 (Eppendorf AG centrifuge 5810R, Germany). Both the macerations and decoctions were carefully
172 filtered using filter paper (Schleicher & Schuell, Ø=150 mm, Germany), and freeze dried for two
173 days in a lyophilizer. Prior to the agar diffusion test, the freeze dried extracts were reconstituted and
174 re-dissolved in their corresponding solvents or in MeOH to a final concentration of 50 mg/mL for
175 the antifungal screening according to the method of Anyanwu and Nwosu, (2014) and Salih et al.,
176 (2017).

177

178 2.5 Fungal strains

179 The fungal strains used in this investigation were obtained from the Division of Pharmaceutical
180 Biosciences, Faculty of Pharmacy, University of Helsinki, Finland. In all, the growth inhibitory
181 activity of the extracts were investigated using five fungal strains (*Candida albicans* ATCC 10231,
182 *Candida glabrata* ATCC 2001, *Candida tropicalis* ATCC 750, *Candida parapsilosis* ATCC 7330,
183 and *Cryptococcus neoformans* ATCC 10226).

184

185 2.6 Antibiotics and pure compounds

186 Amphotericin-B (Sigma-Aldrich, St. Louis MO, USA) and itraconazole (Sigma-Aldrich, St. Louis
187 MO, USA), were used as standard antibiotics for the investigation. Analytical grade piperine and
188 piperlongumine standards ($\geq 97.0\%$ purity) purchased from TCI Europe N.V. (Zwijndrecht,
189 Belgium) were used as standard compounds.

190

191 2.6.1 Agar disk diffusion method

192 Four human pathogenic strains of yeast and *Cryptococcus neoformans*, a yeast-like basidiomycete,
193 were used to investigate the growth inhibitory effects of *P. guineense* extracts. A total of twelve

194 extracts from the fruits and leaf extracts of this plant were tested against the fungi. An agar disk
 195 diffusion method was applied for the initial screening according to Salih et al., (2017). The freeze
 196 dried extracts were re-dissolved in those solvents used originally for their extraction according to
 197 the method of Anyanwu and Nwosu, (2014). Each of the extracts were prepared to a final
 198 concentration of 50 mg/mL (stock solution). Amphotericin-B and itraconazole were used as positive
 199 controls. The antibiotics were dissolved in methanol to a final concentration of 10 mg/mL used for
 200 the test. Sterile petri dishes (\varnothing =15 cm, VWR Finland) were used for the screening. For the
 201 screening, twenty-five mL of sterile base agar (Antibiotic agar No. 2, Difco, VWR Finland) was
 202 applied as a bottom layer into the sterile petri dishes using a sterile, serological pipet (Falcon,
 203 Becton Labware Europe) and allowed to solidify. Thereafter twenty-five mL of Saboraud agar
 204 (OXOID, Thermo Fisher Scientific) was applied as the top layer. The petri dishes were all allowed
 205 to solidify, and then stored in +4°C. The screening started with inoculation of the fungal strains
 206 onto solid Saboraud agar slants which were incubated for 48 h at +37°C. The viable yeast cultures
 207 from the agar slants were used to prepare an inoculum for the test. Fungi from the agar slants were
 208 transferred into 2 mL of 0.9% (w/v) sodium chloride (NaCl) solution in a sterile glass tube using a
 209 sterile inoculation loop. 1 mL of the suspension was transferred into another sterile glass tube, and
 210 the absorbance was measured at 625 nm (UV–Visible Spectrophotometer, Pharmacia LKB-
 211 Biochrom 4060). The other 1 mL of the suspension (sterile part) was diluted with the 0.9% NaCl
 212 solution so that the absorbance at 625 nm becomes 0.1 (this suspension contains approximately 1.5
 213 $\times 10^8$ CFU /mL). 200 μ L of this diluted fungal suspension was spread evenly on each petri dish and
 214 left to dry for some seconds with the lid open. A sterile cork borer (11mm diameter) was used to
 215 make six holes equidistantly from each other on the agar surface of the petri dishes. 200 μ L of the
 216 50 mg/mL plant extracts and 200 μ L of the 10 mg/mL antibiotics were carefully pipetted into the
 217 holes respectively. Methanol, ethanol, hexane and chloroform, 200 μ L of each, were used as solvent
 218 controls respectively. The solvents were found not to inhibit the growth of the fungi. The petri
 219 dishes were incubated at +4 °C for 1 h, and thereafter they were incubated for 48 h at +37 °C. The
 220 diameters of the zones of inhibition were measured with a caliper under a petri dish magnifier and
 221 expressed as the mean of the diameters of three replicates \pm SEM.
 222 The Activity index (AI) of the various extracts were measured in relation to the standard antibiotics
 223 Amphotericin-B and itraconazole according to Fyhrquist et al., (2014).
 224

225 Thus, AI = Inhibition zone of the plant extract

226 Inhibition zone of standard antibiotic

228 2.6.2 Microdilution turbidimetric broth method for MIC and MFC estimation

229 From the result obtained from the agar disk diffusion assay, minimum inhibitory concentration
230 (MIC) was estimated for some selected extracts based on their good antifungal activity. MIC is
231 considered to be the lowest concentration of an extract or compound resulting in the inhibition of at
232 least 90% of the growth of a fungal strain. MIC values were determined using a microdilution
233 turbidimetric broth method based on the guidelines of Clinical and Laboratory Standards Institute
234 (2012). Only extracts which expressed marked antifungal activity in the agar disk diffusion assay
235 were tested for MIC. For the MIC evaluation, two-fold serial dilutions of the extracts from 9.75-
236 2500 µg/mL were prepared in sterile Saboraud broth. Commercial pure compounds, piperine
237 (1 mg/mL concentration in methanol) and piperlongumine (1 mg/mL concentration in methanol)
238 were also two-fold serially diluted in Saboraud broth. Itraconazole and amphotericin-B were each
239 two-fold serially diluted in Saboraud broth from 0.48-125 µg/mL respectively. 96 well microtiter
240 plates were used for the tests (Nunc, Nunclone, Denmark). The fungal cultures were inoculated on
241 Saboraud agar slants or in 5 mL Saboraud broth and grown for 48 hours at +37°C before the test.
242 The absorbance of 1 mL of the 48 h fungal culture was measured for turbidity at $\lambda = 625$ nm using a
243 UV-Visible Spectrophotometer type 1510 (Thermo Fisher Scientific Oy). The absorbance was
244 adjusted to 0.1 at 625 nm (approximately 1.0×10^8 CFU/mL). 100 µL of this suspension $A_{625} = 0.1$
245 was further diluted 100-fold to get a working suspension or inoculum containing 1.0×10^6
246 CFU/mL. 100 µL of this inoculum, and 100 µL of the plant extracts, pure compounds, antibiotics,
247 and solvent controls, were pipetted into the 96 well microtiter plates. Therefore, each well contained
248 5×10^5 CFU/mL. The solvent controls contained a maximum of 5 % (v/v) of each solvent to be
249 tested for toxicity. At this concentration the solvents were found not to be toxic. The growth control
250 (GC wells) contained only the fungal suspension, and the test wells (T wells) contained plant
251 extracts or pure compounds + fungal suspension. Moreover, sample controls wells were prepared
252 for each plant extract/compound to be tested, and these wells contained plant extract/pure
253 compound and the broth only. The microwell plates were incubated for 48 hours in an incubator
254 coupled to a shaker at +37°C. The turbidity of the wells at $\lambda = 620$ nm was recorded using a Victor
255 1420 spectrophotometer (Wallac, Finland). The tests were done in triplicate and the % growth was
256 expressed as the mean of these triplicates \pm standard error of mean (SEM). The minimum fungicidal
257 concentration (MFC) was evaluated by pipetting 100 µL from those wells of the microtiter plate,
258 which contained the MIC concentration as well as 2 and 4 times higher concentrations than their
259 MIC values on petri dishes ($\varnothing = 9$ cm) containing Saboraud agar, and incubating the dishes for 48

hours at +37°C. The MFC was taken as the lowest concentration where no visible growth on the petri dish was observed after the incubation.

3. Results and discussion

3.1 Ethnobotanical survey and field study

An ethnobotanical survey was carried out in Imo state in South-Eastern Nigeria, as shown in Fig. 1. During the interviews and conversations, the various methods of preparations and administrations of the decoctions and concoctions from the roots, leaves, and fruits of *P. guineense* for the treatment of fungal diseases, were widely explained by the traditional healers. 20 traditional healers (14 male and 6 female) were interviewed with the cooperation of their village heads. The participants were between the ages of 40 to 70 years. In the communities visited, it was observed that most of the traditional healers are of older age and have long been practicing traditional medicine. It was also observed that there were more male than female traditional healers and this is in line with previous findings that there are more male than female traditional healers in Africa (Cheikhyoussef, et al., 2011; Ngarivhume et al., 2015). The socio-economic characteristics of the respondents are shown in Table 2.

Table 2. Demographic characteristics of the respondents (n=20)

Parameters	Specifications	Number of Respondents
Gender	Male	14
	Female	6
Occupation	Herbalist	7
	Traditional medical practitioners	5
Age	Traditional healers/ herb sellers	8
	20 - 35	0
	36 - 55	9
	56 - 75	11
Year of practice (experience)	1 – 15 years	6
	16 -30 years	9
	31 years and above	5
Learning/source of knowledge of traditional medicine	Inherited and learnt from parents	12
Level of Education	Training from other herbalists	8
	University education	3
	High school	8
	Primary education	6
	No formal education	3
Religion	Christianity	16
	Traditionalist	4
	Islam	0

Interestingly, most of the traditional healers have remarkable experience in traditional medicine and have inherited and learnt the practice from their parents. All the participants have practiced herbal

281 medicine for a long period of time, and 11 out of the 20 participants have practiced traditional
282 medicine for over 25 years as their only source of income. Notably, some of the participants also
283 had university education and thus some of the information on the preparation of *P. guineense* they
284 also had in written form. One of the traditional healers has a big herbal clinic where he displayed
285 various herbal preparations from *P. guineense* and he is referred to as “Doctor” in his community.
286 It was observed during the course of this study that most of the traditional healers could read and
287 write, that is to say that the trend in traditional medicine in Africa is gradually changing to become
288 a modern profession. It was also observed that some of the traditional healers have books where
289 they recorded the name of each patient and the treatments. From the survey, a variety of herbal
290 preparations from various parts of *P. guineense* were identified. The respondents explained the
291 sources from which they obtain the plant materials for their traditional healing practice. Most of
292 them, apart from traditional healing practice, are also farmers. They have farms within their yards
293 where they planted *P. guineense* and other medicinal plants to be used for the treatment of various
294 infectious diseases and for the purpose of making local profit. The participants explained that they
295 sometimes sell *P. guineense* fruits and leaves in local markets as spices and medicine to the general
296 population. This illustrates that the traditional healers do not have problems in getting the plant
297 materials used for the herbal treatments since they are sold in the local markets. All the respondents
298 gave similar explanations on the symptoms and methods of diagnosis of the fungal infections. The
299 traditional healers explained that they could differentiate between bacterial, viral and fungal
300 infections based on the symptoms they observe in their patients especially when it comes from
301 fungal skin infection and oral thrush. They argued that it was easy for them to identify and
302 differentiate between these infections owing to their long time practice of traditional medicine.
303 They gave some common symptoms for the various fungal infections as shown in Table 3. The
304 method of preparation ranges from hot infusion, decoction in combination with *Xylopi aethiopica*
305 and then the plant material is soaked in mild alcohol as shown Table 3.

306

307 The traditional healers understood the dangers of fungal infections and sometimes send their
308 patients to the government hospitals if the symptoms persist as a result of their failed treatment or
309 lack of proper understanding of the type of fungal infection. The participants explained widely the
310 methods of administration of *P. guineense* extracts. The oral intake is the most common method of
311 administration and these correlates with the findings of Maroyi, (2013) that herbal preparations are
312 mostly administered orally by traditional healers. The traditional healers argued that the decoctions
313 made from the fruits and leaves, prepared in mild alcohol is mostly effective when administered
314 orally for the treatment of fungal diseases expressed by thrush on the tongue or candidal vaginosis.

When asked about the dosage, they further explained that it is usually taken with a small glass tumbler which can be measured to be about 100 mL, 3 to 4 times in a day. It was observed that the leaves and fruits are the most frequently used plant parts for the treatments and this correlates with the findings of Rahmatullah et al., (2012), that the leaves of medicinal plants are used more often in traditional medicine than the other plant parts perhaps because leaves are easy to collect. The traditional healers cooperated with the investigating team and claimed that their methods were effective in the treatment of fungal infections. This study demonstrates that the traditional healers could easily identify *P. guineense* owing to their long practice of the use of extracts of this species in herbal medicine. Our results indicate that *P. guineense* is an important medicinal plant which is highly utilized in the treatment of fungal infections in Imo state, South- Eastern Nigeria and could be a source for the discovery of a new antifungal scaffolds for drug discovery.

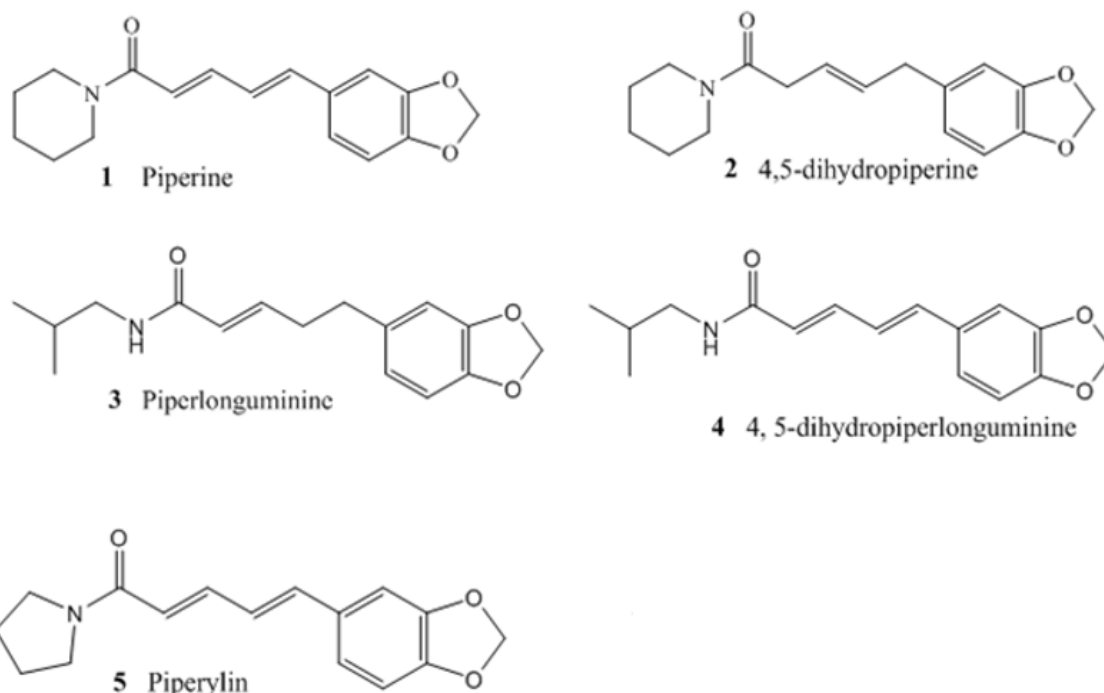
Table 3. List of fungal diseases often treated with *P. guineense* herbal remedies in Imo state, South Eastern Nigeria and the symptoms used by the traditional healers to diagnose the fungal infections. Frequency means the number of healers that mentioned it.

Diseases	Local names in native Igbo language	Symptoms/ Method of diagnosis	Plant parts used	Method of preparation	Administration	Frequency
Skin ringworm	Nwanra, oria ringoringo	Ring-like rashes, and fever	Fruits and leaves	Soaked in alcohol local gin (Kai-kai)	Taken orally	8
Vaginal candidiasis	Oria nwanyi	Vagina itching, unpleasant vaginal discharge, fever	Leaves, fruits, roots	Soaked in alcohol local gin (Kai-kai)	Administered orally	6
Fungal eye infection	Oria anya	Eye itching, swollen of the eye	Leaves	Cold and hot water infusion	Administered in drops in the eye	3
Skin diseases	Oria enuahụ	Rashes, itching, reddish skin, blisters	Fruits and leaves	Cold maceration combined with <i>Zingiber officinale</i>	Applied on the skin	8
Urinary infections	Oria akpaamiri	Painful urination, constant stooling and body weakness	Leaves, and fruits	Soaked in alcohol local gin (Kai-kai)	Administered orally	5
Mouth infections/ Thrush	Obu, Oria Onu, nla	Blisters in the mouth, sour throat, and fever	Fruits and leaves	Decoction combined with <i>Xylopia aethiopica</i>	Used to wash the mouth regularly and taken orally	7
Nail infections/ athlete's foot	Ogbaukwu	Reddish and swollen toes and nails, itchy small blisters	Leaves, and fruits	Soaked in mild alcohol local gin (Kai-kai)	Taken orally	8
Scalp ringworm	Oria Isiakupkpọ, Akpukpọisi	Bald patches, rashes and painful blisters on the head, fever and headache	Fruits and leaves	Soaked in mild alcohol to produce a paste, Hot infusion	Applied as paste on the affected area, infusion taken orally	8

330 The leaf and fruit extracts of *P. guineense* were most frequently used for skin diseases, nail infections/ athlete's foot, scalp ringworm
331 and skin ringworm (recorded 8 times by the 20 traditional healers). Mouth infections/ Thrush and vaginal candidiasis were recorded 7
332 and 6 times respectively.

333

334 From previous reports, *P. guineense* extracts of the seed, fruit and leaf contained piperamide
335 compounds such as piperine, dihydropiperine, piperylin, piperlonguminine,
336 dihydropiperlonguminine, wisanine and dihydrowisanine (Adesina et al, 2003; Scott et al., 2005).
337 However, to the best of our knowledge, no reports are available on the antifungal effects of isolated
338 phytochemicals from *P. guineense*. The chemical structures of piperine, 4, 5-dihydropiperine,
339 piperlonguminine, 4, 5-dihydropiperlonguminine, and piperylin are shown in Figure 2. These
340 bioactive compounds have been identified as the major bioactive compounds in *P. guineense* fruit
341 extract (Scott et al., 2005).



342

343 Fig. 2. Chemical structures of some piperamide compounds in *Piper guineense* fruits

344

345 3.2 Antifungal activity

346 In the present study, a total of 12 extracts of various polarities from the leaf and fruit of *P.*
347 *guineense*, as well as hot water decoctions and cold water macerations were screened for antifungal
348 activity against four yeast strains and *C. neoformans* (Table 4 and 5). Additionally, piperine and
349 piperlongumine commercial piperamide compounds were also screened for antifungal activity. The
350 screening was done based on our ethnobotanical results on the uses of *P. guineense* in West-African

351 traditional medicine for the treatment of fungal diseases (Table 3). For the investigation, some
352 potentially pathogenic fungal strains of *Candida* spp. and *Cryptococcus neoformans*, which could
353 be responsible for some of the fungal diseases treated by the traditional healers were chosen.
354 Among all the fungal strains selected, *C. albicans* is known to be the most significant human
355 pathogenic species of yeast that can cause serious fungal diseases in humans (Brown et al., 2014).
356 The extracts were found to exhibit marked antifungal activity against the various fungi used in our
357 study and MIC values, as low as 39 µg/mL were recorded against some of the fungal strains (Table
358 5). Our results are in agreement with a previous report that fractions, natural compounds and
359 extracts from *P. guineense* can be explored as antifungal agents in the prevention of skin infection
360 (Ngane et al., 2003). Moreover, evaluation of the antifungal potentials of *P. guineense* extracts
361 along with other Cameroonian spices, revealed that the fruit extract of *P. guineense* is effective
362 against *C. albicans*, *C. parapsilosis*, *C. tropicalis*, *C. neoformans*, and *C. glabrata* with MIC values
363 ranging from 1560-6250 µg/mL (Dzoyem et al., 2014). Compared to this study, we recorded even
364 lower MIC and MFC values which could be due to different kinds of extraction methods, Dzoyem
365 et al. (2014) using only methanol: dichloromethane (3:1), but not sequential extraction.
366 Generally, from our study, we observed that the leaf extracts showed a different spectrum of
367 activity compared to the fruits. For example, the hexane leaf extract showed an inhibition zone of
368 21.7 mm against *C. albicans*, whereas the hexane fruit extract only gave an IZ of 14.3 mm against
369 this fungus. The result correlated well with the MIC values, which were recorded to be 78 µg/mL
370 for the leaf extract and 625 µg/mL for the fruit extract (Table 5). A similar trend was observed with
371 the leaf extract in ethanol which gave an inhibition zone of 23.8 mm, whereas the fruit extract gave
372 an IZ of 22.2 mm against *Cryptococcus neoformans*. The MIC values were 78 µg/mL and 156
373 µg/mL, respectively. However, this could be attributed to the fact that the concentration of the
374 antifungal compounds varies between different extracts, thus, this result demonstrates that the
375 leaves might contain a larger quantity of the bioactive compounds responsible for the antifungal
376 activity compared to the fruit extracts. This result could be supported by our ethnobotanical survey
377 results, where the traditional healers explained that the leaves of *P. guineense* are sometimes more
378 active in the treatment of some of the fungal infections and are therefore used more frequently than
379 the fruit for some specific symptoms of fungal infection. Moreover, our results are in accordance
380 with Rahmatullah et al., (2012), who argued that the leaf extracts of medicinal plants are used more
381 often in traditional medicine because they are usually more effective than extracts made from other
382 plant parts.

383

384 Table 4. Antifungal effects of extracts of *P. guineense*, piperine and piperlongumine against four
 385 potentially pathogenic strains of yeast and *Cryptococcus neoformans*. Results obtained with the
 386 agar diffusion method.

Plant extracts and antibiotics	<i>Candida glabrata</i>	<i>Candidaal bicans</i>	<i>Candida parapsilosis</i>	<i>Candida tropicalis</i>	<i>Cryptococcus neoformans</i>
PSMeOH	24,3 ±0,33	20,3 ±0,33	18,0± 0,33	21,3 ± 0,67	19,7 ±0,33
Al. Itra.	1,49	0,80	0,61	0,82	0,66
Al. Amp.	0,71	0,57	0,41	0,73	0,52
PSCHCL ₃	23,3 ± 0,33	17,7±0,33	14,7± 0,67	16,2±.0,17	18,2±0,17
Al. Itra.	1,43	0,70	0,49	0,62	0,61
Al. Amp.	0,68	0,50	0,34	0,55	0,48
PSHex	20,3 ±0,33	14,3± 0,33	15,3± 0,67	11,0±0,58	14,3±0,33
Al. Itra.	1,24	0,57	0,52	0,42	0,48
Al. Amp.	0,59	0,40	0,35	0,38	0,38
PSH ₂ O*	NA	NA	NA	NA	NA
PSHH ₂ O	NA	NA	NA	NA	NA
PSEthanol	NT	21,8 ± 0,17	NT	19,8 ± 0,17	22,2 ±0,17
Al. Itra.	NT	0,86	NT	0,76	0,75
Al. Amp.	NT	0,61	NT	0,68	0,58
PLMeOH	20,7 ± 0,31	19,7 ±0,33	28,7 ±0,67	17,3± 0,33	21,3 ±0,67
Al. Itra.	1,27	0,78	0,97	0,66	0,72
Al. Amp.	0,60	0,55	0,66	0,59	0,56
PLCHCL ₃	11,7± 0,33	18,0±0,00	13,7±.0,17	15,5± 0,50	17,3±0,33
Al. Itra.	0,71	0,71	0,46	0,59	0,58
Al. Amp.	0,34	0,50	0,31	0,53	0,46
PLHex	13,7± 0,33	21,7 ± 0,33	17,8±0,17	12,0± 0,58	11,7± 0,33
Al. Itra.	0,84	0,86	0,60	0,46	0,39
Al. Amp.	NT	0,61	0,41	0,41	0,31
PLH ₂ O*	NA	NA	NA	NA	NA
PLHH ₂ O	NA	NA	NA	NA	NA
PLEthanol	NT	19,8 ± 0,17	NT	18,2±0,17	23,8 ±0,17
Al. Itra.	NT	0,78	NT	0,69	0,80
Al. Amp	NT	0,56	NT	0,62	0,63
Pure compounds					
Piperine	17,3± 0,67	18,3±.0,33	23,3 ± 0,33	16,7±0,67	11,0±.0,00
Al. Itra.	1,06	0,72	0,79	0,64	0,37
Al. Amp	0,50	0,51	0,53	0,57	0,29
Piperlongumine	14,7± 0,33	19,3 ±.0,33	16,3± 0,33	18,0±0,00	11,8±0,17
Al. Itra.	0,90	0,76	0,55	0,69	0,40
Al. Amp	0,43	0,54	0,37	0,61	0,31
Antifungal drugs					
Itraconazole	16,3± 0,33	25,3 ±.0,33	29,7 ± 0,67	26,2 ±0,17	29,7 ±0,33
Al. Amp	0,48	0,71	0,68	0,89	0,78
Amphotericin B	34,3 ± 0,33	35,7 ± 0,33	43,7 ± 0,67	29,3 ±.0,67	38,0 ± 0,00
Al. Itra.	2,10	1,41	1,47	1,12	1,28
Solvent controls					
Methanol	2,3	2,8	NA	NA	NA
Chloroform	NA	NA	NA	NA	NA
Hexane	3,8	2,3	NA	NA	NA
Ethanol	NT	3, 7	NT	2, 3	NA

387 The diameter of the zones of inhibition in mm as the mean of triplicates (n = 3) ± SEM (standard error of mean). 200µL of extracts
 388 and fractions (50 mg/mL), as well as pure compounds and antibiotics (10 mg/mL) were applied in the wells. PSMethOH, *P. guineense*
 389 fruit methanol extract; PSCHCL₃, *P. guineense* fruit chloroform extract; PSHex, *P. guineense* fruit hexane extract; PSH₂O*, *P.*
 390 *guineense* fruit cold water maceration; PSHH₂O, *P. guineense* fruit hot water decoction; PSEthanol, *P. guineense* fruit ethanol
 391 extract; PLMeOH, *P. guineense* leaf methanol extract; PLCHCL₃, *P. guineense* leaf chloroform extract; PLHex, *P. guineense* leaf
 392 hexane extract; PLH₂O*, *P. guineense* leaf cold water maceration; PLHH₂O, *P. guineense* leaf hot water decoction; PLEthanol, *P.*

393 *guineense* leaf ethanol extract; AI Itra., activity index in relation to itraconazole; AI Amp., activity index in relation to amphotericin
394 B; NA, not active and NT, not tested. The results marked with bold show promising antifungal activity.

395 Interestingly, the water extracts were not active against the fungal strains evaluated in our study.
396 Our results are consistent with a previous report that cold water macerations and hot water
397 decoctions of *P. guineense* are not active against *C. albicans* and other *Candida* spp. (Okigbo and
398 Igwe, 2007). Contrary to our results, water extracts of *P. guineense* are used in traditional medicine
399 for fungal infections, even though it is sometimes administered as herbal formulations in
400 combination with other medicinal plants, and often mixed with the local gin (Table 3). In our study,
401 we observed that some of the alcohol and n-hexane extracts have marked antifungal activity with
402 MIC values as low as 39 µg/mL, and the inhibitory activity recorded with these extracts could
403 result from their piperamide compounds, which are not well dissolved in water. It is possible that
404 the water extracts contain only small quantities of piperamide alkaloids, which might explain their
405 inactivity against the yeasts. The water extracts may contain other bioactive compounds than
406 alkaloids, and thus *P. guineense* water extracts might be better to use in combination with other
407 medicinal plants as herbal remedy for treatment of fungal infections in traditional medicine, which
408 could explain the uses of *P. guineense* water extracts as concoctions for fungal infections in West-
409 African traditional medicine.

410 Our results demonstrate that the ethanol fruit and leaf extracts of *P. guineense* give good growth
411 inhibition against *C. albicans* (IZ 21.8 mm and 19.8 mm, respectively) (Table 4), and accordingly
412 both extracts gave low MIC values of 78 µg/mL against *C. albicans*. *C. albicans* is an opportunistic
413 pathogenic fungus that can cause serious systemic infections in humans, affecting mostly
414 individuals with compromised immune defences, such as HIV/AIDS patients and elderly people
415 (Brown et al., 2014). *C. albicans* is the main causative agent of candidiasis and our results are in
416 agreement with the traditional uses of the fruits and leaves soaked in mild alcohol for the treatment
417 of vaginal candidiasis (Table 3). Our result is also consistent with a previous report by Ejele et al.,
418 (2012), that ethanol extracts of *P. guineense* have significant activity against *C. albicans*. We also
419 observed significant and promising growth inhibitory results of 20.3 mm and 19.7 mm with the
420 methanol fruit and leaf extracts against *C. albicans*. Moreover, the chloroform extracts were active
421 against *C. albicans* with a MIC value of 78 µg/mL for the leaf and fruit extracts. In our screenings,
422 piperlongumine was found to be very active against *C. albicans* with a MIC value of 39 µg/mL,
423 while piperine gave a MIC of 78 µg/mL. To the best of our knowledge, the inhibitory efficacy of
424 piperlongumine has not been tested against *C. albicans*, and little information is available on the
425 activity of piperine on *Candida* species. Since there is a report of multi-drug resistance in *C.*
426 *albicans* (Gulshan and Moye-Rowley, (2007), our result demonstrates that piperlongumine and

427 piperine could be scaffolds for new natural plant derived antifungal agents to combat multi-drug
 428 resistance in *Candida* strains. More research should be conducted to ascertain the antifungal
 429 mechanism of action of piperamide alkaloids as well as on producing extracts of *P. guineense*
 430 leaves or fruits which would be standardized to their piperamide alkaloid contents.
 431 In comparison to our results with the plant extracts and alkaloids, our amphotericin-B standard drug
 432 showed a large inhibitory zone of 35.7 mm and a MIC of 0.48 µg/mL against *C. albicans*. The
 433 activity of *P. guineense* extracts and its piperamide alkaloids on *C. albicans* demonstrates that its
 434 extracts and alkaloids could be effectively utilized in combinations with conventional antifungals
 435 for the treatment of infections caused by antibiotic resistant *C. albicans* strains.
 436 We also recorded promising growth inhibitory effects against *C. glabrata*, the second most
 437 significant human pathogenic species of yeast. The methanol, chloroform, and hexane extracts were
 438 all more active against *C. glabrata* than itraconazole (Table 4). Methanol and chloroform extracts
 439 gave the largest inhibition zones against *C. glabrata* (IZ ranging from 24.3 mm to 20.7 mm) (Table
 440 4). The MIC value for the chloroform fruit extract was 78 µg/mL. Notably, itraconazole was only
 441 moderately active against *C. glabrata*, with an inhibition zone of 16.3mm and a MIC value of 15.6
 442 µg/mL. Drug-resistance against itraconazole has been found to be associated with 5% of
 443 nosocomial isolates of *C. glabrata* (Walker et al., 2013), and our results demonstrate that extracts of
 444 *P. guineense* could be sources for new antifungal scaffolds and drug adjuvants against drug resistant
 445 *C. glabrata*. A very low MIC value of 0.48 µg/mL was recorded with amphotericin B against *C.*
 446 *glabrata*, and our result is in agreement with the more frequent use of amphotericin B for the
 447 treatment of severe infections caused by *C. glabrata* (Mario et al., 2012). Piperine was active
 448 against *C. glabrata* with a MIC of 78 µg/mL, while piperlongumine was moderately active with a
 449 MIC value of 156 µg/mL. Apart from *C. albicans*, *C. glabrata* is another important fungal pathogen
 450 responsible for systemic infections which are associated with high mortality rate (Pfaller et al.,
 451 2004; 2007). Our results demonstrate that the extracts and the piperamide compounds could be
 452 possible sources and antifungal drug scaffolds for the discovery of a new antifungal drugs and drug
 453 adjuvants for the treatment of systemic infections caused by *C. glabrata*.

454
 455 Table 5. Minimum inhibitory concentration and minimum fungicidal concentrations (MIC and
 456 MFC) in µg/mL of fruit and leaf extracts of *Piper guineense* and the alkaloids piperine and
 457 piperlongumine

Plant extracts and antibiotics	<i>Candida</i> <i>glabrata</i>	<i>Candida</i> <i>albicans</i>	<i>Candida</i> <i>parapsilosis</i>	<i>Candida</i> <i>tropicalis</i>	<i>Cryptococcus</i> <i>neoformans</i>
-----------------------------------	-----------------------------------	-----------------------------------	---------------------------------------	-------------------------------------	--

PSMeOH					
MIC	39	39	625	39	156
MFC	78	78	1250	78	312
PSCHCl ₃					
MIC	78	78	625	312	156
MFC	312	156	125	625	312
PSHex					
MIC	78	625	625	625	1250
MFC	1250	1250	1250	1250	2500
PSEthanol					
MIC	NT	78	NT	78	156
MFC	NT	156	NT	156	625
PLMeOH					
MIC	78	156	312	78	156
MFC	156	312	625	156	312
PLCHCl ₃					
MIC	78	78	625	312	156
MFC	156	156	1250	625	312
PLHex					
MIC	312	78	625	312	1250
MFC	625	625	1250	625	2500
PLEthanol					
MIC	NT	78	NT	156	78
MFC	NT	156	NT	312	156
Piperine					
MIC	78	78	39	78	312
MFC	156	156	78	156	625
Piperlongumine					
MIC	156	39	156	78	312
MFC	312	78	312	156	625
Amphotericin B					
MIC	0.48	0.48	0.48	0.48	0.97
Itraconazole					
MIC	0.97	15.6	0.97	0.48	0.48

459 MIC value represents mean of triplicates. MIC, minimum inhibitory concentration; MFC, minimum fungicidal concentration. For
460 key to other abbreviations, see Table 4. Only extracts with promising antifungal effect in the agar disk diffusion assay were tested for
461 MIC. The MFC was evaluated by taking 100 µL from the wells containing extracts of the MIC concentration and 2 times and 4 times
462 the MIC concentrations and inoculating on a petri dish for 48 hours. The MFC was taken as the lowest concentration where no
463 visible growth was observed after the incubation. The results marked with bold show promising MIC values.

464 For *Candida tropicalis*, the methanol fruit extract gave the largest inhibition zone (IZ 21.3 mm) and
465 a MIC value of 39 µg/mL. The MIC value for piperine against this fungus was 78 µg/mL and thus
466 was less effective than the methanol fruit extract. Piperine has been reported to be one of the main
467 piperamide compounds found in *P. guineense* (Scott et al., 2005). Our result could indicate that
468 some other piperamide compounds in the methanol fruit extract could be responsible for the good
469 growth inhibitory effects of the extract, or then that these compounds act in synergy together with
470 piperine to produce a lower MIC than that of pure piperine. The other extracts of the fruits were
471 moderately active against *C. tropicalis* with inhibition zones ranging from 19.8 to 11.0 mm (Table

472 4). The methanol leaf extract was also active against the fungus with a MIC of 78 µg/mL. The MIC
473 value of the ethanol fruit and leaf extracts was found to be 78 µg/mL for both. In contrast to the
474 other extracts, the decoctions and macerations showed no growth inhibition against *C. tropicalis*.
475 All extracts, except for the macerations and decoctions were active against *C. parapsilopsis* with
476 MIC values ranging from 312 to 625 µg/mL. The largest inhibition zone of 28.7 mm and 18.0 mm
477 were recorded for a methanol leaf and fruit extract, respectively. A mixture of methanol-
478 dichloromethane (3:1 v/v) of the fruit extract of *P. guineense* has been previously found to be active
479 against this fungus with a MIC value of 625 µg/mL (Dzoyem et al., 2014). Our result indicates that
480 pure methanol is a better extractant for antifungals than methanol-dichloromethane (3:1), since we
481 recorded a larger inhibition zone of 28.7 mm and a lower MIC value of 312 µg/mL with the
482 methanol leaf extract compared to Dzoyem et al., (2014). The growth inhibitory effect of the
483 ethanol extracts were not tested against *C. parapsilopsis* due to lack of material. Piperine and
484 piperlongumine were active against *C. parapsilopsis* with inhibition zones of 23.3 and 16.3 mm,
485 respectively. The MIC values for piperine and piperlongumine were found to be 39 µg/mL and 156
486 µg/mL, respectively. Thus, piperine could be responsible for some of the good activity displayed by
487 the methanol leaf extract.

488

489 In our study, *P. guineense* extracts were active against *Cryptococcus neoformans*. The largest
490 inhibitory activity was observed with the ethanol leaf and fruit extracts (23.8 mm and 22.2 mm)
491 (Table 4). The MIC values for the ethanol leaf and fruit extracts were in agreement with the agar
492 diffusion result (78 µg/mL and 156 µg/mL respectively). The methanol leaf and fruit extracts
493 recorded inhibition zones of 21.3 mm and 19.7 mm respectively, and their MIC values were
494 recorded to be 156 µg/mL for both. It was observed that the leaf extracts of *P. guineense* were more
495 active against this fungus than the fruits extracts. The chloroform fruit and leaf extracts were active
496 against the fungus with inhibition zones of 18.2 mm and 17.3 mm. The hexane fruit and leaf
497 extracts were moderately active also (IZ 14.3 mm and 11.7 mm). The MIC values for the
498 chloroform and hexane extracts against this fungus were 156 µg/mL and 2500 µg/mL, respectively.
499 Our result is consistent with one previous report that *P. guineense* fruit extract has growth inhibitory
500 activity against *C. neoformans* with a MIC of 312 µg/mL (Dzoyem et al., 2014). Piperine and
501 piperlongumine were active with a MIC of 312 µg/mL for both.

502

503 **Conclusion**

504 *P. guineense* extracts, piperine and piperlongumine have been found to possess interesting
505 antifungal properties as recorded in this study. The antifungal efficacy of the methanol, ethanol,

chloroform and *n*-hexane extracts of the fruit and leaf of *P. guineense*, as well as piperine and piperlongumine on various *C. albicans* and non-*albicans Candida* strains supports the use of *P. guineense* in the treatment of fungal infections in traditional medicine and demonstrates that extracts from this plant could contain bioactive compounds that could be utilized as therapeutic agents or scaffolds for the production of new antifungal drugs to treat diseases associated with *Candida* species and *Cryptococcus neoformans*. The study demonstrated that the extracts and piperamide compounds exert promising antifungal properties against pathogenic *Candida albicans* and other non-*albicans Candida* strains. Moreover, various extracts, especially from *P. guineense* leaves, showed a promising growth inhibitory profile against *Cryptococcus neoformans* which is known to cause life-threatening meningitis in immunocompromised individuals. In view of this current knowledge on alkaloids of *P. guineense* as antifungal agents, we recommend that additional research should be done to evaluate the *in vivo* antifungal properties of the extracts and piperamide compounds from *P. guineense* with some animal models. The marked inhibitory activity recorded with piperine and piperlongumine against *C. albicans* revealed that they could be good antifungal agents. Further research is needed to evaluate the inhibitory activity of these piperamide compounds in combination with currently known standard antibiotics because piperine and piperlongumine could possibly enhance the effects of the currently available antifungal drugs for the treatment of fungal infections. This study has clarified and broadened the traditional knowledge on the uses of *P. guineense* as an antifungal remedy in African traditional medicine. Systematic studies are in progress and for further investigations, HPLC-DAD and UHPLC/Q-TOF MS analysis are currently being conducted by our research team to characterize and identify the compounds present in the extracts with promising antifungal activity.

528

529 **Author's contributions**

530 E.E.M, Y.H, P.F, H.V, and C.A designed the experiment.

531 E.E.M and C.A conducted the field work, ethnobotanical survey and documentations in Nigeria.

532 E.E.M, Y.H, H.V and P.F performed the *in vitro* antifungal screening and MIC estimations. The

533 manuscript was first drafted by E.E.M, C.A, and P.F. The manuscript was further revised and

534 corrected by H.V, Y.H and P.F. All the authors have read, agreed and approved the final version of the manuscript.

536

537 **Conflict of interest**

538 The authors indicate no potential conflicts of interest.

539

540 **Ethical issues**

541 The ethical consideration of the project was done appropriately according to the authors' institution
542 and was approved by the institutional ethics committee. The ethnobotanical survey was done with
543 the approval of the village heads and according to the national institutional rules of the indigenous
544 people.

545

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552 healers and herb sellers for giving their time.

553

554 **Appendix A. Supplementary material**

555 Questionnaire used for the ethnobotanical survey on the traditional use of *Piper guineense* for the
556 treatment of fungal infections in Imo state, South Eastern Nigeria.

557 We are researchers from the University of Helsinki, Finland and Federal University of Technology
558 Owerri, Imo state, Nigeria. We would like to know the various methods that you use for utilization
559 of Uziza (*Piper guineense*) for the treatment of fungal infections.

560 1. Gender Male ☐ Female ☐

561 2. What is your name (optional).....

562 3. What is your major occupation? : a). Herbalist ☐ b). Traditional medical practitioner ☐

563 c). Traditional healer/ herb seller ☐ d). Other ☐

564 4. What is your age? : a). 20-35 ☐ 36 – 55 ☐ 56 – 75 ☐

565 5. For how long have you been practicing traditional medicine? : a) 1 – 15 years ☐

566 16 -30 years ☐ 31 years and above ☐

567 6. What is your source of knowledge of traditional medicine? : a). Inherited/learnt from parents ☐

568 Training from other herbalists ☐ Other ☐

569 7. What is your level of education? : a). University education ☐ High school ☐

570 Primary education ☐ No formal education ☐

571 8. What is your religion? : a) Christianity ☐ Traditionalist ☐ Islam ☐ Other ☐

572 9. Can you name some of the fungal diseases that you treat with Uziza (*Piper guineense*)?

.....
.....
.....

10. How often do you treat these diseases?

.....

11. What are the symptoms and how do you diagnose this fungal infections?

Fungal infections	Local name	Method of diagnosis/ symptoms

12. Which part of the plant do you often use?: Fruit ☐ Leaves ☐ Roots ☐ Other ☐

13. How do you get the *Piper guineense* used for the treatment? : a) Cultivate them ☐ b) Buy from the market ☐ c) Collect from the wild forest and bushes ☐ d) Other ☐

14. What are the method of preparations and administrations?

Fungal diseases	Method of preparation	Mode of administration

573 15. What are the dosage of administration and how long does the treatment last?

574
 575 16. What do you do if you observe that the patient is not responding to your antifungal treatment?
 576
 577 Do you have some general comment about the possible challenges you face in your daily treatment
 578 of fungal infections using *Piper guineense* extracts?
 579

580
 581 Thank you for your cooperation.
 582

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